

General Assembly

Raised Bill No. 5430

February Session, 2022

LCO No. 3086



Referred to Committee on PUBLIC HEALTH

Introduced by: (PH)

AN ACT CONCERNING OPIOIDS.

Be it enacted by the Senate and House of Representatives in General Assembly convened:

- 1 Section 1. Section 20-14s of the general statutes is repealed and the
- 2 following is substituted in lieu thereof (*Effective July 1, 2022*):

3 A prescribing practitioner, as defined in section 20-14c, who

4 prescribes an opioid drug, as defined in section 20-14o, for the treatment

5 of pain for a patient for a duration greater than twelve weeks shall

6 establish a treatment agreement with the patient or discuss a care plan

7 for the chronic use of opioids with the patient. The treatment agreement

8 or care plan shall, at a minimum, include treatment goals, risks of using

opioids, urine drug screens and expectations regarding the continuing

10 treatment of pain with opioids, such as situations requiring

11 discontinuation of opioid treatment and, to the extent possible,

12 nonopioid treatment options, including, but not limited to

13 manipulation, chiropractic, spinal cord stimulation, massage therapy,

14 acupuncture, physical therapy and other treatment regimens or

modalities. A record of the treatment agreement or care plan shall be

16 recorded in the patient's medical record.

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Sec. 2. Subdivision (20) of section 21a-240 of the 2022 supplement to the general statutes is repealed and the following is substituted in lieu thereof (*Effective July 1, 2022*):

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(20) (A) "Drug paraphernalia" [refers to] means equipment, products and materials of any kind [which] that are used, intended for use or designed for use in planting, propagating, cultivating, growing, harvesting, manufacturing, compounding, converting, producing, processing, preparing, testing, analyzing, packaging, repackaging, storing, containing or concealing, or ingesting, inhaling or otherwise introducing into the human body, any controlled substance contrary to the provisions of this chapter including, but not limited to: (i) Kits intended for use or designed for use in planting, propagating, cultivating, growing or harvesting of any species of plant [which] that is a controlled substance or from which a controlled substance can be derived; (ii) kits used, intended for use or designed for use in manufacturing, compounding, converting, producing, processing or preparing controlled substances; (iii) isomerization devices used [,] or intended for use in increasing the potency of any species of plant [which] that is a controlled substance; (iv) testing equipment used, intended for use or designed for use in identifying or analyzing the strength, effectiveness or purity of controlled substances; (v) dilutents and adulterants, [such as] including, but not limited to, quinine hydrochloride, mannitol, mannite, dextrose and lactose used, intended for use or designed for use in cutting controlled substances; (vi) separation gins and sifters used, intended for use or designed for use in removing twigs and seeds from, or in otherwise cleaning or refining, marijuana; (vii) capsules and other containers used, intended for use or designed for use in packaging small quantities of controlled substances; (viii) containers and other objects used, intended for use or designed for use in storing or concealing controlled substances; (ix) objects used, intended for use or designed for use in ingesting, inhaling, or otherwise introducing marijuana, cocaine, hashish, or hashish oil into the human body, [such as: Metal,] including, but not limited to, wooden, acrylic, glass, stone, plastic or ceramic pipes with screens, permanent screens,

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hashish heads or punctured metal bowls; water pipes; carburetion tubes 51 52 and devices; smoking and carburetion masks; roach clips; [: Meaning 53 objects used to hold burning material, such as a marijuana cigarette, that 54 has become too small or too short to be held in the hand; miniature 55 cocaine spoons [,] and cocaine vials; chamber pipes; carburetor pipes; 56 electric pipes; air-driven pipes; chillums; bongs; [or] ice pipes [or] and 57 chillers. "Drug paraphernalia" does not include a product used by a 58 manufacturer licensed pursuant to this chapter for the activities 59 permitted under the license or by an individual to test any substance 60 prior to injection, inhalation or ingestion of the substance to prevent 61 accidental overdose by injection, inhalation or ingestion of the 62 substance, provided the licensed manufacturer or individual is not 63 using the product to engage in the unlicensed manufacturing or distribution of controlled substances. As used in this subdivision, "roach 64 65 clip" means an object used to hold burning material, including, but not 66 limited to, a marijuana cigarette, that has become too small or too short 67 to be held between the fingers;

(B) "Factory" means any place used for the manufacturing, mixing, compounding, refining, processing, packaging, distributing, storing, keeping, holding, administering or assembling illegal substances contrary to the provisions of this chapter, or any building, rooms or location which contains equipment or paraphernalia used for this purpose;

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- Sec. 3. Section 20-633c of the general statutes is repealed and the following is substituted in lieu thereof (*Effective July 1, 2022*):
 - (a) A person who is licensed as a pharmacist under part II of this chapter [and is certified in accordance with subsection (b) of this section] may prescribe, in good faith, an opioid antagonist, as defined in section 17a-714a. Such pharmacist shall (1) provide appropriate training regarding the administration of such opioid antagonist to the person to whom the opioid antagonist is dispensed, and (2) maintain a record of such dispensing and the training required pursuant to this chapter.

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[(b) A pharmacist may only prescribe an opioid antagonist pursuant to this section if the pharmacist has been trained and certified by a program approved by the Commissioner of Consumer Protection.]

- [(c)] (b) A pharmacist who prescribes an opioid antagonist in compliance with this section shall be deemed not to have violated any standard of care for a pharmacist.
- [(d) The provisions of this section shall apply only to a pharmacist certified in accordance with subsection (b) of this section.] (c) No pharmacist may delegate or direct any other person to prescribe an opioid antagonist or train any person in the administration of such opioid antagonist pursuant to the provisions of subsection (a) of this section.
- [(e)] (d) The Commissioner of Consumer Protection may adopt regulations, in accordance with chapter 54, to implement the provisions of this section.
- 98 Sec. 4. Section 20-633d of the general statutes is repealed and the following is substituted in lieu thereof (*Effective July 1, 2022*):
 - (a) A prescribing practitioner, as defined in section 20-14c, who is authorized to prescribe an opioid antagonist, as defined in section 17a-714a, and a pharmacy may enter into an agreement for a medical protocol standing order at such pharmacy allowing a pharmacist licensed under part II of this chapter to dispense an opioid antagonist that is (1) administered by an intranasal application delivery system or an auto-injection delivery system, (2) approved by the federal Food and Drug Administration, and (3) dispensed to any person at risk of experiencing an overdose of an opioid drug, as defined in 42 CFR 8.2, or to a family member, friend or other person in a position to assist a person at risk of experiencing an overdose of an opioid drug.
 - (b) Any such medical protocol standing order shall be deemed issued for a legitimate medical purpose in the usual course of the prescribing practitioner's professional practice. The pharmacy shall provide the

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- 114 Department of Consumer Protection with a copy of every medical
- 115 protocol standing order agreement entered into with a prescribing
- 116 practitioner under this section.
- [(c) A pharmacist may only dispense an opioid antagonist pursuant
- to a medical protocol standing order if the pharmacist has been trained
- and certified as part of a program approved by the Commissioner of
- 120 Consumer Protection.]
- [(d)] (c) A pharmacist who dispenses an opioid antagonist pursuant
- to a medical protocol standing order shall (1) provide appropriate
- training regarding the administration of such opioid antagonist to the
- 124 person to whom the opioid antagonist is dispensed, (2) maintain a
- 125 record of such dispensing and the training required pursuant to this
- 126 chapter, and (3) send a copy of the record of such dispensing to the
- 127 prescribing practitioner who entered into an agreement for a medical
- 128 protocol standing order with the pharmacy.
- [(e)] (d) A pharmacist who dispenses an opioid antagonist in
- accordance with the provisions of this section shall be deemed not to
- 131 have violated any standard of care for a pharmacist.
- [(f)] (e) The commissioner may adopt regulations, in accordance with
- chapter 54, to implement the provisions of this section.
- Sec. 5. Section 21a-286 of the general statutes is repealed and the
- following is substituted in lieu thereof (*Effective July 1, 2022*):
- 136 (a) For purposes of this section:
- 137 (1) "Opioid antagonist" shall have the meaning set forth in section
- 138 17a-714a.
- 139 (2) "Prescribing practitioner" shall have the meaning set forth in
- 140 section 20-14c.
- 141 (3) "Pharmacist" shall have the meaning set forth in section 20-609a.

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(b) A prescribing practitioner or a pharmacist [certified to prescribe] who prescribes naloxone pursuant to section 20-633c, as amended by this act, may enter into an agreement with a law enforcement agency, emergency medical service provider, government agency or community health organization related to the distribution and administration of an opioid antagonist for the reversal of an opioid overdose. The prescribing practitioner or pharmacist shall provide training to persons who will distribute or administer the opioid antagonist pursuant to the terms of the agreement. Persons other than the prescribing practitioner or pharmacist shall receive training in the distribution or administration of opioid antagonists prior to distributing or administering an opioid antagonist. The agreement shall address the storage, handling, labeling, recalls and recordkeeping of opioid antagonists by the law enforcement agency, emergency medical service provider, government agency or community health organization which is party to the agreement.

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- (c) A prescribing practitioner or pharmacist who enters into an agreement pursuant to subsection (b) of this section shall not be liable for damages in a civil action or subject to administrative or criminal prosecution for the administration or dispensing of an opioid antagonist by such law enforcement agency, emergency medical service provider, government agency or community health organization.
- 163 The Commissioner of Consumer Protection may adopt regulations, in accordance with the provisions of chapter 54, to implement the provisions of this section.
- 166 Sec. 6. Subsection (a) of section 21a-252 of the general statutes is 167 repealed and the following is substituted in lieu thereof (Effective from 168 passage):
 - (a) A physician, in good faith and in the course of the physician's professional practice only, may prescribe, administer and dispense controlled substances, or may cause the same to be administered by a physician assistant, nurse or intern under the physician's direction and supervision, for demonstrable physical or mental disorders but not for

LCO No. 3086 6 of 11 drug dependence except in accordance with state and federal laws and regulations adopted thereunder. Notwithstanding the provisions of this subsection the Department of Consumer Protection may approve protocols allowing the dispensing of take-home doses of methadone, by a registered nurse or licensed practical nurse, to outpatients in duly licensed substance [abuse] use disorder treatment facilities, including, but not limited to, requesting approval of an exception, through the Department of Mental Health and Addiction Services pursuant to subsection (d) of section 17a-450, to the unsupervised take-home medication requirements set forth in 42 CFR 8.12(i), as amended from time to time, that are necessary to (1) dispense up to twenty-eight days of take-home doses of methadone to a stable patient if, in the professional medical judgment of the registered nurse or licensed practical nurse, such nurse believes the patient can safely tolerate such amount of methadone, and (2) dispense up to fourteen days of takehome doses of methadone to a less stable patient if, in the professional medical judgment of the registered nurse or licensed practical nurse, such nurse believes the patient can safely tolerate such amount of methadone. Such dispensing shall be done pursuant to the order of a licensed prescribing practitioner and using computerized dispensing equipment into which bulk supplies of methadone are dispensed by a pharmacist. The quantity of methadone dispensed by such nurse shall not exceed at any one time that amount allowed under federal or state statutes or regulations governing the treatment of drug dependent patients unless the Department of Mental Health and Addiction Services has approved an exception from the relevant federal statutes or regulations to such allowed amount pursuant to subsection (d) of section 17a-450. The Department of Consumer Protection shall conduct inspections of such treatment facilities to ensure that the computerized dispensing equipment and related dispensing procedures documented in the approved protocols are adhered to.

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Sec. 7. (*Effective from passage*) Not later than January 1, 2023, the Commissioner of Mental Health and Addiction Services shall report, in accordance with the provisions of section 11-4a of the general statutes,

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208 to the joint standing committee of the General Assembly having 209 cognizance of matters relating to public health regarding the sober 210 living homes that have reported their certified status to the Department 211 of Mental Health and Addiction Services pursuant to section 17a-716 of 212 the general statutes. Such report shall include, but not be limited to, the 213 following information: (1) The existence, administration and success of 214 any voucher program that provides financial assistance to individuals, 215 including residents of a sober living home, to alleviate the financial 216 burden of obtaining substance use disorder services, including 217 addiction treatment and rehabilitation services; (2) whether any 218 additional funding is necessary to support residents of sober living 219 homes in obtaining substance use disorder services; and (3) 220 recommendations for any legislative changes necessary to support 221 residents of sober living homes in obtaining substance use disorder 222 services.

Sec. 8. Section 21a-317 of the general statutes is repealed and the following is substituted in lieu thereof (*Effective July 1, 2022*):

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Every practitioner who distributes, administers or dispenses any controlled substance or who proposes to engage in distributing, prescribing, administering or dispensing any controlled substance within this state shall (1) obtain a certificate of registration issued by the Commissioner of Consumer Protection in accordance with the provisions of this chapter, [and] (2) if the practitioner is engaged in prescribing a controlled substance, register for access to the electronic prescription drug monitoring program established pursuant to subsection (j) of section 21a-254 [. Registration for access to said program shall be in a manner prescribed by said commissioner.] in a manner prescribed by the commissioner, and (3) if the practitioner is engaged in transporting a controlled substance for the purpose of treating a patient in a location that is different than the address that the practitioner provided to the Department of Consumer Protection as a registrant, as defined in section 21a-240, as amended by this act, notify the department, in a manner prescribed by the commissioner, of the intent to transport such controlled substance and, after dispensing such

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controlled substance, return any remaining amount of such controlled substance to a secure location at the address provided to the department. If the practitioner cannot return any remaining amount of such controlled substance to such address, the commissioner may approve an alternate location, provided such location is also approved by the federal Drug Enforcement Agency, or any successor agency. The practitioner shall report any dispensation by the practitioner of a controlled substance that occurs at a location other than the address provided to the department to the prescription drug monitoring program pursuant to subsection (j) of section 21a-254 upon returning to such address.

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Sec. 9. (NEW) (*Effective July 1, 2022*) On or before January 1, 2023, and quarterly thereafter, the Departments of Mental Health and Addiction Services, Consumer Protection, Social Services and Correction shall share with the Department of Public Health data collected in the normal course of said departments' business in addressing the opioid epidemic in the state. The Commissioner of Public Health shall collect, organize and analyze such data, along with the same type of data collected by the Department of Public Health, to (1) establish metrics for prescribing opioid drugs, as defined in section 20-140 of the general statutes, and treating persons with opioid use disorder or persons who have experienced an overdose of an opioid drug, and (2) update such metrics on a quarterly basis, if deemed necessary by the commissioner.

Sec. 10. Subsection (j) of section 17a-451 of the general statutes is repealed and the following is substituted in lieu thereof (*Effective July 1*, 2022):

(j) The commissioner shall be responsible for developing and implementing the Connecticut comprehensive plan for prevention, treatment and reduction of alcohol and drug abuse problems to be known as the state substance abuse plan. Such plan shall include a mission statement, a vision statement and goals for providing treatment and recovery support services to adults with substance use disorders. The plan shall be developed by July 1, 2010, and thereafter shall be

triennially updated by July first of the respective year. The commissioner shall develop such plan, mission statement, a vision statement and goals after consultation with: (1) The Connecticut Alcohol and Drug Policy Council established pursuant to section 17a-667; (2) the Criminal Justice Policy Advisory Commission established pursuant to section 18-87j; (3) the subregional planning and action councils established pursuant to section 17a-671; (4) clients and their families, including those involved with the criminal justice system; (5) treatment providers; and (6) other interested stakeholders. The plan shall outline the action steps, time frames and resources needed to meet specified goals and shall, at a minimum, address: (A) Access to services, both prior to and following admission to treatment; (B) the provision of comprehensive assessments to those requesting treatment, including individuals with co-occurring conditions; (C) quality of treatment services and promotion of research-based and evidence-based best practices and models; (D) an appropriate array of prevention, treatment and recovery services along with a sustained continuum of care; (E) outcome measures of specific treatment and recovery services in the overall system of care; (F) information regarding the status of treatment program availability for pregnant women, including statistical and demographic data concerning pregnant women and women with children in treatment and on waiting lists for treatment; (G) department policies and guidelines concerning recovery-oriented care; (H) provisions of the community reentry strategy concerning substance abuse treatment and recovery services needed by the offender population as developed by the Criminal Justice Policy and Planning Division within the Office of Policy and Management; (I) an evaluation of the Connecticut Alcohol and Drug Policy Council's plan described in section 17a-667 and any recommendations for changes to such plan; [and] (J) a summary of data maintained in the central repository, described in subsection (o) of this section; and (K) department policies, guidelines and practices aimed at reducing the negative personal and public health impacts of behavior associated with alcohol and drug abuse, including, but not limited to, the abuse of an opioid drug, as defined in section 20-14o. The plan shall define measures and set

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benchmarks for the overall treatment system and for each state-operated program. Measures and benchmarks specified in the plan shall include, but not be limited to, the time required to receive substance abuse assessments and treatment services either from state agencies directly or through the private provider network funded by state agencies, the percentage of clients who should receive a treatment episode of ninety days or greater, treatment provision rates with respect to those requesting treatment, connection to the appropriate level of care rates, treatment completion rates and treatment success rates as measured by improved client outcomes in the areas of substance use, employment, housing and involvement with the criminal justice system.

This act shall take effect as follows and shall amend the following sections:		
Section 1	July 1, 2022	20-14s
Sec. 2	July 1, 2022	21a-240(20)
Sec. 3	July 1, 2022	20-633c
Sec. 4	July 1, 2022	20-633d
Sec. 5	July 1, 2022	21a-286
Sec. 6	from passage	21a-252(a)
Sec. 7	from passage	New section
Sec. 8	July 1, 2022	21a-317
Sec. 9	July 1, 2022	New section
Sec. 10	July 1, 2022	17a-451(j)

Statement of Purpose:

To combat the opioid epidemic in the state.

[Proposed deletions are enclosed in brackets. Proposed additions are indicated by underline, except that when the entire text of a bill or resolution or a section of a bill or resolution is new, it is not underlined.]

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